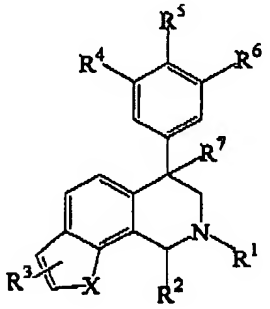
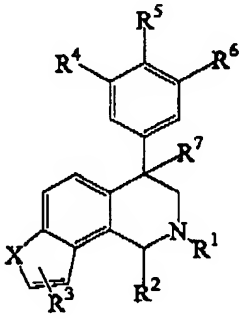
Amendments to the Claims:

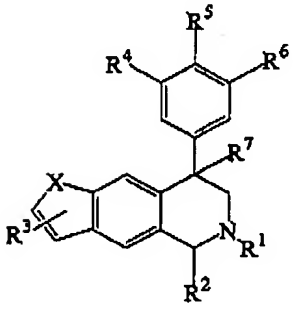
1. (Currently Amended) A method of treating ~~chronic or neuropathic pain, treating or preventing migraine headache, or treating~~ urge, stress or mixed urinary incontinence comprising administration of an effective amount of a compound selected from one of the Formulae IA, IB, IIA, IIB, IIIA or IIIB



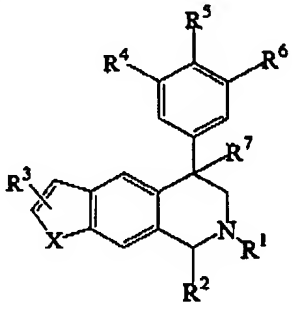
IA



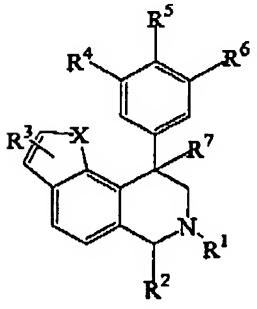
IB



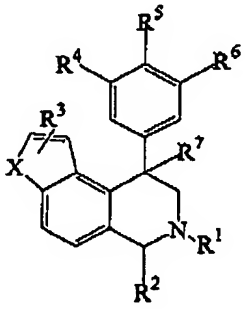
IIA



IIB



IIIA



IIIB

wherein:

R^1 is selected from the group consisting of C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl, C_4-C_7 cycloalkylalkyl and benzyl, each of which is optionally substituted with 1 to 3 substituents independently selected at each occurrence from C_1-C_3 alkyl, halogen, -CN, $-OR^8$ and $-NR^8R^9$;

R^2 is selected from the group consisting of H, C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl, C_4-C_7 cycloalkylalkyl and C_1-C_6 haloalkyl;

R^3 is selected from the group consisting of H, halogen, C_1-C_6 alkyl, C_1-C_6 haloalkyl and C_3-C_6 cycloalkyl, wherein C_1-C_6 alkyl, C_1-C_6 haloalkyl and C_3-C_6 cycloalkyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from OR^8 and NR^8R^9;

R^4, R^5, and R^6 are each independently selected at each occurrence thereof from the group consisting of H, halogen, $-OR^{10}$, $-NO_2$, $-NR^{10}R^{11}$, $-NR^{10}C(O)R^{11}$, $-NR^{10}C(O)NR^{11}R^{12}$, $-S(O)_nR^{11}$, -CN, $-C(O)R^{11}$, $-C(O)_2R^{11}$, $-C(O)NR^{11}R^{12}$, C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl and C_4-C_7 cycloalkylalkyl, wherein each of C_1-C_6 alkyl, C_2-C_6 alkenyl, C_2-C_6 alkynyl, C_3-C_6 cycloalkyl and C_4-C_7 cycloalkylalkyl are optionally substituted with 1 to 3 substituents independently selected at each occurrence with from C_1-C_3 alkyl, halogen, =0, -CN, $-OR^8$, $-NR^8R^9$ and phenyl, and wherein phenyl is optionally substituted 1-3 substituents selected independently at each occurrence from halogen, -CN, C_1-C_4 alkyl, C_1-C_4 haloalkyl, $-OR^8$ and $-NR^8R^9$;

alternatively R^5 and R^6 taken together are $-O-C(R^{11})_2-O-$;

R^7 is selected from the group consisting of H, halogen and OR^{10};

R^8 and R^9 are each independently selected from the group consisting of H, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxyalkyl, C_1-C_4 alkoxyalkylalkyl, C_3-C_6 cycloalkyl, C_4-C_7 cyclooalkylalkyl, $-C(O)R^{12}$, phenyl and benzyl, wherein phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from halogen, cyano, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxy and C_1-C_4 haloalkoxy, or R^8 and R^9 are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine ring;

R^{10} is selected from the group consisting of H, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxyalkyl, C_3-C_6 cycloalkyl, C_4-C_7 cycloalkylalkyl, -C(O)R^{12}, phenyl and benzyl, wherein phenyl and benzyl are optionally substituted with 1 to 3 substituents selected. independently at each occurrence from halogen, -NH_2, -OH, cyano, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxy and C_1-C_4 haloalkoxy;

R^{11} is selected from the group consisting of H, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxyalkyl, C_3-C_6 cycloalkyl, C_4-C_7 cycloalkylalkyl, phenyl and benzyl, where phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from halogen, -NH_2, -OH, cyano, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxy and C_1-C_4 haloalkoxy, or R^{10} and R^{11} are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, N-methylpiperazine, morpholine, or thiomorpholine ring, with the proviso that only one of R^8 and R9 or R^{10} and R^{11} are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, piperaine, N-methylpiperazine, morpholine, or thiomorpholine ring;

R^{12} is selected from the group consisting of C_1-C_4 alkyl, C_1-C_4 haloalkyl and phenyl;

X is selected from the group consisting of 0, NR^{13} and S;

the ring containing X is selected from furan, pyrrole, thiophene, dihydrofuran, dihydropyrrole, and dihydrothiophene; n is 0, 1, or 2; and,

R^{13} is selected from the group consisting of H, C_1-C_6 alkyl, benzyl and phenyl, wherein C_1-C_6 alkyl, benzyl and phenyl are optionally substituted with 1-3 substituents independently at each occurrence from halogen, -NH_2, -OH, cyano, C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_1-C_4 alkoxy and C_1-C_4 haloalkoxy;

or a pharmaceutically acceptable salt thereof or an isomer or prodrug thereof to a patient in need thereof.

2. (Original) A method of claim 1, wherein R^1 is C_1-C_6 alkyl.

3. (Original) A method of claim 2, wherein R^1 is CH_3.

4. (Original) A method of claim 1, wherein R^2 is H, C_1-C_6 alkyl, C_3-C_6 cycloalkyl, or C_1-C_6 haloalkyl.

5. (Original) A method of claim 4, wherein R^2 is H or C_1-C_6 alkyl.

6. (Original) A method of claim 5, wherein R^2 is H.

7. (Original) A method of claim 1, wherein R^3 is at each occurrence thereof independently H, halogen, C_1-C_6 alkyl, or C_1-C_6 alkyl substituted with from 1 to 3 of OR^8 or NR^8R^9.

8. (Original) A method of claim 7, wherein R^3 is H or C_1-C_6 alkyl.

9. (Original) A method of claim 8, wherein R^3 is H.

10. (Original) A method of claim 1, wherein R^1 is CH_3, R^2 is H and R^3 is H.

11. (Original) A method of claim 1, wherein R^4, R^5 and R^6 are each independently H, halogen, C_1-C_6 alkyl or -OR^{10}.

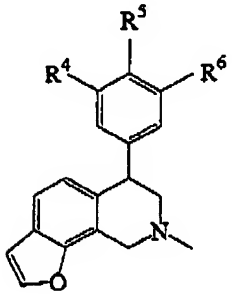
12. (Original) A method of claim 11, wherein at least one of R^4, R^5 and R^6 is H.

13. (Original) A method of claim 12, wherein each of R^4, R^5 and R^6 are H.

14. (Original) A method of claim 12, wherein one of R^4, R^5 and R^6 is halogen.

15. (Original) A method of claim 1, wherein R^1 is CH_3, R^2 and R^3 are each H, and at least one of R^4, R^5, and R^6 is H.

16. (Original) A method of claim 1 wherein the compound is a compound of Formula (10):



(10)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

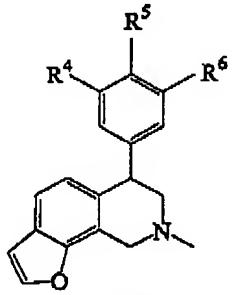
a compound of Formula (10) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (10) wherein R^4 is H, R^5 is Me and R^6 is H;

a compound of Formula (10) wherein R^4 is Cl, R^5 is H and R^6 is H; and

a compound of Formula (10) wherein R^4 is H, R^5 is F and R^6 is H.

17. (Original) A method of claim 1 wherein the compound is a compound of Formula (20):



(20)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (20) wherein R^4 is H, R^5 is H and R^6 is H;

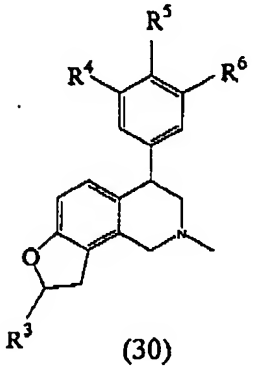
a compound of Formula (20) wherein R^4 is H, R^5 is Me and R^6 is H;

a compound of Formula (20) wherein R^4 is H, R^5 is Cl and R^6 is H;

a compound of Formula (20) wherein R^4 is H, R^5 is F and R^6 is H; and

a compound of Formula (20) wherein R^4 is F, R^5 is H and R^6 is F.

18. (Original) A method of claim 1 wherein the compound is a compound of Formula (30):

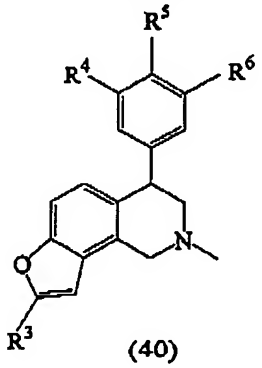


(30)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is H and R^6 is H;
a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is F and R^6 is H;
a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is F;
a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is H;
a compound of Formula (30) wherein R^3 is H, R^4 is Cl, R^5 is H and R^6 is H;
a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is H;
a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is F;
a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is Cl;
a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is Cl;
a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is OMe and R^6 is H; and
a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is H.

19. (Original) A method of claim 1 wherein the compound is a compound of Formula (40):

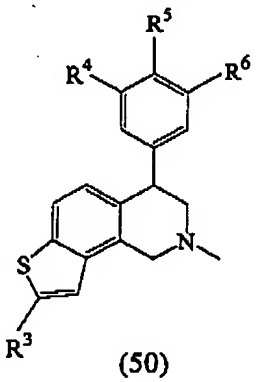


(40)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is H and R^6 is H;
a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is F and R^6 is H;
a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is F;
a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is H;
a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is H;
a compound of Formula (40) wherein R^3 is H, R^4 is Cl, R^5 is H and R^6 is H;
a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is H;
a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is F;
a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is Cl;
a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is Cl;
a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is 0Me and R^6 is H;
a compound of Formula (40) wherein R^3 is Me, R^4 is H, R^5 is H and R^6 is H;
a compound of Formula (40) wherein R^3 is Et, R^4 is H, R^5 is H and R^6 is H; and
a compound of Formula (40) wherein R^3 is CH_20H, R^4 is H, R^5 is H and R^6 is H.

20. (Original) A method of claim 1 wherein the compound is a compound of Formula (50):

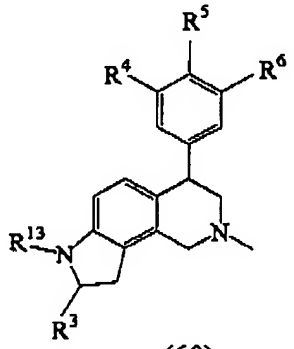


(50)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (50) wherein R^3 is H, R^4 is H, R^5 is H and R^6 is H.

21. (Original) A method of claim 1 wherein the compound is a compound of Formula (60):



(60)

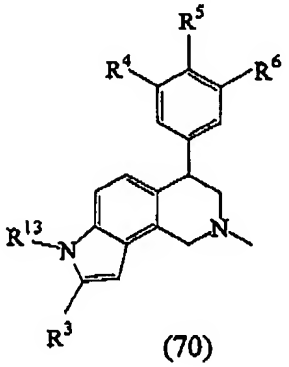
or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is H;
a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Me;
a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Et;
a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is H;
a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is Me;
a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is F and R^{13} is H;
a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is F and R^{13} is Me;
a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is H;
a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is Me;
a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is H and R^{13} is H;
a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is H and R^{13} is H;
a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is H;
a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is Me;
a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is H;

and

a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is Me.

22. (Original) A method of claim 1 wherein the compound is a compound of Formula (70):



(70)

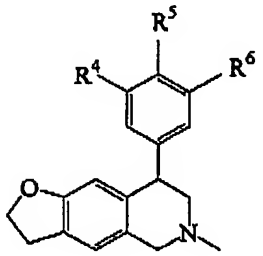
or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is H;
a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Me;
a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Et;
a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Bn;
a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is H;
a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is Me;
a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is F and R^{13} is Me;
a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is H;
a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is Me;
a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is H and R^{13} is H;
a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is H and R^{13} is Me;
a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is H and R^{13} is H;
a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is H;
a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is Me;
a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is H;

and

a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is Me.

23. (Original) A method of claim 1 wherein the compound is a compound of Formula (80):



(80)

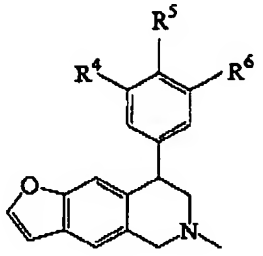
or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (80) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (80) wherein R^4 is H, R^5 is F and R^6 is H; and

a compound of Formula (80) wherein R^4 is H, R^5 is F and R^6 is F.

24. (Original) A method of claim 1 wherein the compound is a compound of Formula (90):



(90)

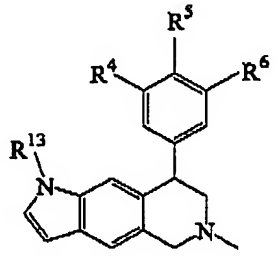
or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (90) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (90) wherein R^4 is H, R^5 is F and R^6 is F; and

a compound of Formula (90) wherein R^4 is H, R^5 is F and R^6 is H.

25. (Original) A method of claim 1 wherein the compound is a compound of Formula (100):

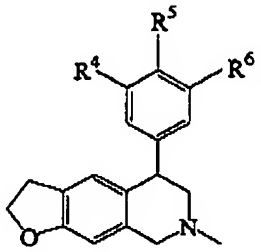


(100)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (100) wherein R^4 is H, R^5 is H, R^6 is H and R^{13} is H.

26. (Original) A method of claim 1 wherein the compound is a compound of Formula (110):



(110)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (110) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is F;

a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is H;

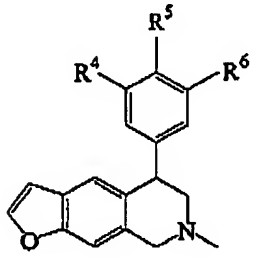
a compound of Formula (110) wherein R^4 is H, R^5 is H and R^6 is Cl;

a compound of Formula (110) wherein R^4 is H, R^5 is Cl and R^6 is F;

a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is Cl; and

a compound of Formula (110) wherein R^4 is H, R^5 is 0Me and R^6 is H.

27. (Original) A method of claim 1 wherein the compound is a compound of Formula (120):



(120)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (120) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (120) wherein R^4 is H, R^5 is F and R^6 is F;

a compound of Formula (120) wherein R^4 is H, R^5 is F and R^6 is H;

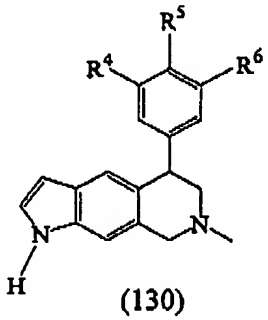
a compound of Formula (120) wherein R^4 is H, R^5 is H and R^6 is Cl;

a compound of Formula (120) wherein R^4 is H, R^5 is Cl and R^6 is F;

a compound of Formula (120) wherein R^4 is H, R^5 is 0Me and R^6 is H; and

a compound of Formula (120) wherein R^4 is H, R^5 is F and R^6 is Cl.

28. (Original) A method of claim 1 wherein the compound is a compound of Formula (130):



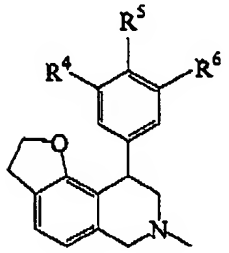
(130)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (130) wherein R^4 is H, R^5 is H and R^6 is H; and

a compound of Formula (130) wherein R^4 is H, R^5 is Bn and R^6 is H.

29. (Original) A method of claim 1 wherein the compound is a compound of Formula (140):



(140)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (140) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (140) wherein R^4 is H, R^5 is F and R^6 is H;

a compound of Formula (140) wherein R^4 is H, R^5 is F and R^6 is Cl;

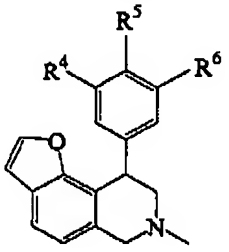
a compound of Formula (140) wherein R^4 is H, R^5 is Cl and R^6 is F;

a compound of Formula (140) wherein R^4 is H, R^5 is H and R^6 is Cl;

a compound of Formula (140) wherein R^4 is H, R^5 is 0Me and R^6 is H;

a compound of Formula (140) wherein R^4 is H, R^5 is F and R^6 is F.

30. (Original) A method of claim 1 wherein the compound is a compound of Formula (150):



(150)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (150) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (150) wherein R^4 is H, R^5 is F and R^6 is H;

a compound of Formula (150) wherein R^4 is H, R^5 is F and R^6 is Cl;

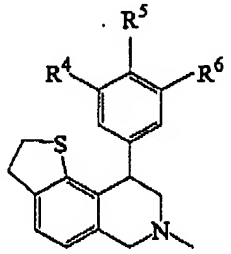
a compound of Formula (150) wherein R^4 is H, R^5 is Cl and R^6 is F;

a compound of Formula (150) wherein R^4 is H, R^5 is H and R^6 is Cl;

a compound of Formula (150) wherein R^4 is H, R^5 is 0Me and R^6 is H; and

a compound of Formula (150) wherein R^4 is H, R^5 is F and R^6 is F.

31. (Original) A method of claim 1 wherein the compound is a compound of Formula (160):

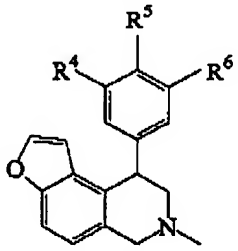


(160)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (160) wherein R^4 is H, R^5 is H and R^6 is H.

32. (Original) A method of claim 1 wherein the compound is a compound of Formula (170):



(170)

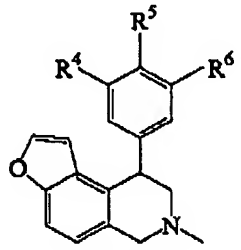
or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (170) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (170) wherein R^4 is H, R^5 is F and R^6 is H; and

a compound of Formula (170) wherein R^4 is H, R^5 is F and R^6 is F.

33. (Original) A method of claim 1 wherein the compound is a compound of Formula (180):



(180)

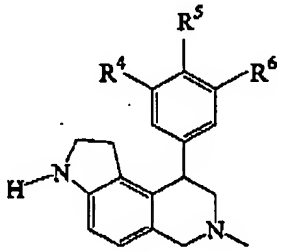
or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (180) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (180) wherein R^4 is H, R^5 is F and R^6 is H; and

a compound of Formula (180) wherein R^4 is H, R^5 is F and R^6 is F.

34. (Original) A method of claim 1 wherein the compound is a compound of Formula (190):

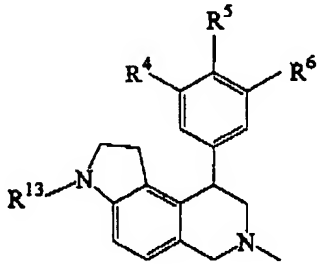


(190)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (190) wherein R^4 is H, R^5 is H and R^6 is H.

35. (Original) A method of claim 1 wherein the compound is a compound of Formula (200):



(200)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (200) wherein R^4 is H, R^5 is H, R^6 is H and R^{13} is H; and

a compound of Formula (200) wherein R^4 is H, R^5 is H, R^6 is H and R^{13} is Me.

36. (Original) A method of claim 1 wherein the compound is selected from the group consisting of:

(R)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2,3-h]isoquinoline;

(S)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2, 3-h]isoquinoline;

(R)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3,2-g]isoquinoline;

(S)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;

(R)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;

(S)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;

(R)-4-(3,4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2, 3-h]isoquinoline;

(S)-4-(3, 4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;

(R)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2, 3-h]isoquinoline;

(S)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2, 3-h]isoquinoline;

(R)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4- tetrahydro-furo[2, 3-h]isoquinoline;

(S)-4-(4-chloro-phenyl)-2- methyl-1,2,3,4-tetrahydrofuro[2,3- h]isoquinoline;

(R)-8-methyl- 6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;

(S)-8-methyl-6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;

(R)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;

(S)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(R)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3, 4-tetrahydrofuro[2,3- h]isoquinoline;

(S)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(R)-2-methyl-4-phenyl2,3,4,7-tetrahydro-1H-pyrrolo[2,3-h]isoquinoline; and

(S)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H-pyrrolo[2, 3-h]isoquinoline.

37. (Original) A method of claim 1 wherein the compound is selected from the group consisting of:

(+)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2, 3-h]isoquinoline;

(-)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2,3-h]isoquinoline;

(+)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;

(-)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;

(+)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(-)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydro-furo[2,3-h]isoquinoline;

(+)-4-(3,4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(-)-4-(3,4- difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(+)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(-)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(+)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;

(-)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(+)-8-methyl-6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2h]isoquinoline;

(-)-8-methyl- 6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;

(+)-4-(4- fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(-)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;

(+)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(-)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(+)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H-pyrrolo[2,3-h]isoquinoline; and

(-)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H- pyrrolo[2,3-h]isoquinoline.